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Classification Data
NEWS 5 FEB 02 Simultaneous left and right truncation (SLART) added
for CERAB, COMPUAB, ELCOM, and SOLIDSTATE
NEWS 6 FEB 02 GENBANK enhanced with SET PLURALS and SET SPELLING
NEWS 7 FEB 06 Patent sequence location (PSL) data added to USGENE
NEWS 8 FEB 10 COMPENDEX reloaded and enhanced
NEWS 9 FEB 11 WTEXTILES reloaded and enhanced
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NEWS 16 FEB 25 USGENE enhanced with patent family and legal status
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NEWS 19 MAR 11 ESBIOBASE reloaded and enhanced
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for nanomaterial substances
NEWS 21 MAR 23 CA/CAPLUS enhanced with more than 250,000 patent
equivalents from China
NEWS 22 MAR 30 IMSPATENTS reloaded and enhanced
NEWS 23 APR 03 CAS coverage of exemplified prophetic substances
enhanced

NEWS EXPRESS JUNE 27 08 CURRENT WINDOWS VERSION IS V8.3,
AND CURRENT DISCOVER FILE IS DATED 23 JUNE 2008.

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* * * * * STN Columbus * * * * *

FILE 'HOME' ENTERED AT 15:39:41 ON 07 APR 2009

=> fil reg		
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	ENTRY	SESSION
FULL ESTIMATED COST	0.22	0.22

FILE 'REGISTRY' ENTERED AT 15:39:51 ON 07 APR 2009
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STRUCTURE FILE UPDATES: 6 APR 2009 HIGHEST RN 1132745-38-0
DICTIONARY FILE UPDATES: 6 APR 2009 HIGHEST RN 1132745-38-0

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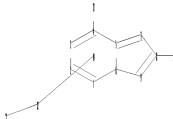
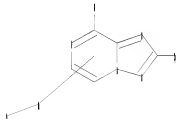
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Uploading C:\Program Files\STNEXP\Queries\10552305newest.str



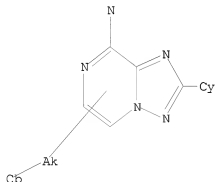
chain nodes :
11 12 13 15
ring nodes :
1 2 3 4 5 6 7 8 9
chain bonds :

4-11 8-15 12-13
 ring bonds :
 1-2 1-6 2-3 3-4 4-5 5-6 5-7 6-9 7-8 8-9
 exact/norm bonds :
 1-2 1-6 2-3 3-4 4-5 4-11 5-6 5-7 6-9 7-8 8-9 8-15 12-13
 isolated ring systems :
 containing 1 :

Match level :
 1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 6:Atom 7:Atom 8:Atom 9:Atom 11:CLASS
 12:CLASS 13:Atom 14:CLASS 15:Atom
 Generic attributes :
 13:
 Saturation : Saturated
 Type of Ring System : Monocyclic

L1 STRUCTURE UPLOADED

=> d l1
 L1 HAS NO ANSWERS
 L1 STR



Structure attributes must be viewed using STN Express query preparation.

=> s l1
 SAMPLE SEARCH INITIATED 15:40:07 FILE 'REGISTRY'
 SAMPLE SCREEN SEARCH COMPLETED - 543 TO ITERATE
 100.0% PROCESSED 543 ITERATIONS 2 ANSWERS
 SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE **COMPLETE**
 BATCH **COMPLETE**
 PROJECTED ITERATIONS: 9462 TO 12258
 PROJECTED ANSWERS: 2 TO 124

L2 2 SEA SSS SAM L1

=> s l1 sss full
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 FULL SCREEN SEARCH COMPLETED - 10474 TO ITERATE

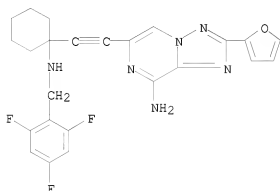
100.0% PROCESSED 10474 ITERATIONS
SEARCH TIME: 00.00.01

10 ANSWERS

L3 10 SEA \$\$\$ FUL L1

=> d scan

L3 10 ANSWERS REGISTRY COPYRIGHT 2009 ACS on STN
IN [1,2,4]Triazolo[1,5-a]pyrazin-8-amine,
2-(2-furanyl)-6-[2-[1-[(2,4,6-
trifluorophenyl)methyl]amino]cyclohexyl]ethynyl]-
MF C24 H21 F3 N6 O



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

HOW MANY MORE ANSWERS DO YOU WISH TO SCAN? (1):0

=> fil cap

COST IN U.S. DOLLARS

SINCE FILE

TOTAL

ENTRY

SESSION

FULL ESTIMATED COST

185.88

186.10

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FILE COVERS 1907 - 7 Apr 2009 VOL 150 ISS 15

FILE LAST UPDATED: 6 Apr 2009 (20090406/ED)

Caplus now includes complete International Patent Classification (IPC) reclassification data for the third quarter of 2008.

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This file contains CAS Registry Numbers for easy and accurate substance identification.

=> d his

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FILE 'REGISTRY' ENTERED AT 15:39:51 ON 07 APR 2009

L1 STRUCTURE UPLOADED

L2 2 S L1

L3 10 S L1 SSS FULL

FILE 'CAPLUS' ENTERED AT 15:40:22 ON 07 APR 2009

=> s l3

L4 2 L3

=> d 1-2 ibib abs hitstr

L4 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2009 ACS on STN

ACCESSION NUMBER: 2005:74649 CAPLUS

DOCUMENT NUMBER: 142:298062

TITLE: Synthesis of alkyne derivatives of a novel triazolopyrazine as A2A adenosine receptor antagonists
AUTHOR(S): Yao, Gang; Haque, Serajul; Sha, Li; Kumaravel, Gnanasambandam; Wang, Joy; Engber, Thomas M.; Whalley, Eric T.; Conlon, Patrick R.; Chang, Hexi; Kiesman, William F.; Petter, Russell C.

CORPORATE SOURCE: Departments of Medicinal Chemistry and Pharmacology, Biogen Idec, Cambridge, MA, 02142, USA

SOURCE: Bioorganic & Medicinal Chemistry Letters (2005), 15(3), 511-515

CODEN: BMCLE8; ISSN: 0960-894X

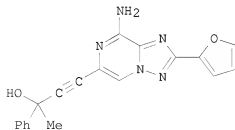
PUBLISHER: Elsevier B.V.

DOCUMENT TYPE: Journal

LANGUAGE: English

OTHER SOURCE(S): CASREACT 142:298062

GI



I

AB A [1,2,4]triazolo[1,5-a]pyrazine core was synthesized and coupled with terminal acetylenes. The structure-activity relationship of the alkynes, e.g., 1, from this template was studied for their in vitro and in vivo

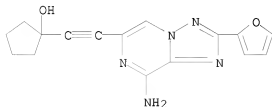
adenosine A2A receptor antagonism. Selected compds. from this series were shown to have potent in vitro and in vivo activities against adenosine A2A receptor. I was found to be orally active at 3 mg/kg in both a mouse catalepsy model and a 6-hydroxydopamine-lesioned rat model.

IT 785049-27-6P 785049-29-8P

RL: PAC (Pharmacological activity); PRP (Properties); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation)
(preparation, A2A adenosine receptor affinity, Parkinson's disease efficacy, and SAR of alkynyltriazolopyridazines via Sonogashira coupling of amino(furanyl)bromotriazolopyridazine with alkynes)

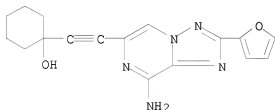
RN 785049-27-6 CAPLUS

CN Cyclopentanol, 1-[2-[8-amino-2-(2-furanyl)[1,2,4]triazolo[1,5-a]pyrazin-6-yl]ethynyl]- (CA INDEX NAME)



RN 785049-29-8 CAPLUS

CN Cyclohexanol, 1-[2-[8-amino-2-(2-furanyl)[1,2,4]triazolo[1,5-a]pyrazin-6-yl]ethynyl]- (CA INDEX NAME)

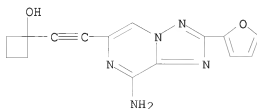


IT 785049-26-5P 785049-37-8P 785049-51-6P

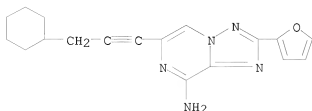
RL: PAC (Pharmacological activity); PRP (Properties); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation)
(preparation, A2A adenosine receptor affinity, and SAR of alkynyltriazolopyridazines via amination of aminodibromopyrazine with carbamate followed by condensation with furancarboxaldehyde, cyclization, and Sonogashira coupling with alkynes)

RN 785049-26-5 CAPLUS

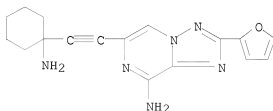
CN Cyclobutanol, 1-[2-[8-amino-2-(2-furanyl)[1,2,4]triazolo[1,5-a]pyrazin-6-yl]ethynyl]- (CA INDEX NAME)



RN 785049-37-8 CAPLUS
 CN [1,2,4]Triazolo[1,5-a]pyrazin-8-amine,
 6-(3-cyclohexyl-1-propyn-1-yl)-2-(2-furanyl)- (CA INDEX NAME)



RN 785049-51-6 CAPLUS
 CN [1,2,4]Triazolo[1,5-a]pyrazin-8-amine,
 6-[2-(1-aminocyclohexyl)ethynyl]-2-(2-furanyl)- (CA INDEX NAME)



REFERENCE COUNT: 29 THERE ARE 29 CITED REFERENCES AVAILABLE FOR THIS
 RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 2 OF 2 CAPLUS COPYRIGHT 2009 ACS on STN

ACCESSION NUMBER: 2004:902386 CAPLUS

DOCUMENT NUMBER: 141:395583

TITLE: Preparation of triazolopyrazines as A2a adenosine
 receptor antagonists for the treatment of Parkinson's
 disease

INVENTOR(S): Dowling, James; Yao, Gang; Chang, Hexi; Peng, Hairuo;
 Vessels, Jeffrey; Petter, Russell C.; Kumaravel,
 Gnanasambandam

PATENT ASSIGNEE(S): Biogen Idec Ma Inc., USA

SOURCE: PCT Int. Appl., 100 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

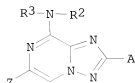
LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

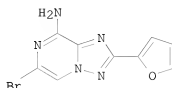
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2004092177	A1	20041028	WO 2004-US11006	20040409
W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW			
RW:	BW, GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE, SI,			

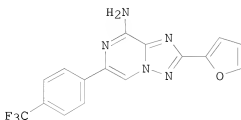
SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN,
 TD, TG
 EP 1615931 A1 20060118 EP 2004-759356 20040409
 R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
 IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, PL, SK, HR
 US 20070010520 A1 20070111 US 2006-552305 20060829
 PRIORITY APPLN. INFO.: US 2003-461546P P 20030409
 WO 2004-US11006 W 20040409
 OTHER SOURCE(S): CASREACT 141:395583; MARPAT 141:395583
 GI



I



II

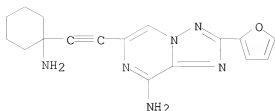


III

AB Title compds. I [A = aryl, heteroaryl; R₂, R₃ = H, alkyl, cycloalkyl, etc.; Z = -X₁-L-X₂-Y-X₃-R₁; X₁, X₂, X₃ = bond, alkylene, alkenylene, etc.; L = bond or cyclic-linker] and their pharmaceutically acceptable salts and N-oxides were prepared For example, coupling of 4-trifluoromethylphenylboronic acid and bromophenyl II, e.g., prepared from furan-2-carbonitrile in 3-steps, afforded claimed triazolopyrazine III. In A_{2a} adenosine receptor binding assays, compds. I exhibited K_i values less than 10 μM. Compds. I are claimed useful for the treatment of Parkinson's disease.

IT 785049-51-6P
 RL: PAC (Pharmacological activity); RCT (Reactant); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); RACT (Reactant or reagent); USES (Uses)
 (preparation of triazolopyrazines as A_{2a} adenosine receptor antagonists for the treatment of Parkinson's disease)

RN 785049-51-6 CAPLUS
 CN [1,2,4]Triazolo[1,5-a]pyrazin-8-amine,
 6-[2-(1-aminocyclohexyl)ethynyl]-2-(2-furanyl)- (CA INDEX NAME)



IT 785049-20-9P 785049-26-5P 785049-27-6P

785049-28-7P 785049-29-8P 785049-34-5P

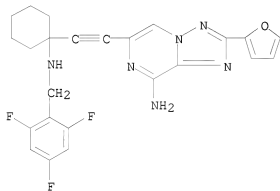
785049-35-6P 785049-37-8P 785049-38-9P

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(preparation of triazolopyrazines as A2a adenosine receptor antagonists for the treatment of Parkinson's disease)

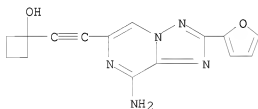
RN 785049-20-9 CAPLUS

CN [1,2,4]Triazolo[1,5-a]pyrazin-8-amine, 2-(2-furanyl)-6-[2-[1-[(2,4,6-trifluorophenyl)methylamino]cyclohexyl]ethynyl]- (CA INDEX NAME)



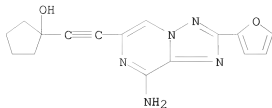
RN 785049-26-5 CAPLUS

CN Cyclobutanol, 1-[2-[8-amino-2-(2-furanyl)[1,2,4]triazolo[1,5-a]pyrazin-6-yl]ethynyl]- (CA INDEX NAME)



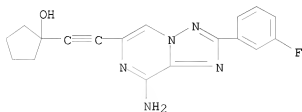
RN 785049-27-6 CAPLUS

CN Cyclopentanol, 1-[2-[8-amino-2-(2-furanyl)[1,2,4]triazolo[1,5-a]pyrazin-6-yl]ethynyl]- (CA INDEX NAME)



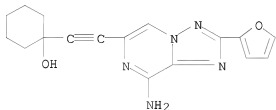
RN 785049-28-7 CAPLUS

CN Cyclopentanol, 1-[2-[8-amino-2-(3-fluorophenyl)[1,2,4]triazolo[1,5-a]pyrazin-6-yl]ethynyl]- (CA INDEX NAME)



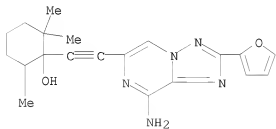
RN 785049-29-8 CAPLUS

CN Cyclohexanol, 1-[2-[8-amino-2-(2-furanyl)[1,2,4]triazolo[1,5-a]pyrazin-6-yl]ethynyl]- (CA INDEX NAME)



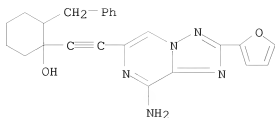
RN 785049-34-5 CAPLUS

CN Cyclohexanol, 1-[2-[8-amino-2-(2-furanyl)[1,2,4]triazolo[1,5-a]pyrazin-6-yl]ethynyl]-2,2,6-trimethyl- (CA INDEX NAME)

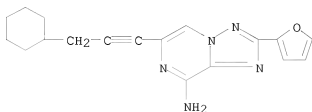


RN 785049-35-6 CAPLUS

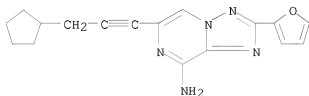
CN Cyclohexanol, 1-[2-[8-amino-2-(2-furanyl)[1,2,4]triazolo[1,5-a]pyrazin-6-yl]ethynyl]-2-(phenylmethyl)- (CA INDEX NAME)



RN 785049-37-8 CAPLUS
 CN [1,2,4]Triazolo[1,5-a]pyrazin-8-amine,
 6-(3-cyclohexyl-1-propyn-1-yl)-2-(2-furanyl)- (CA INDEX NAME)



RN 785049-38-9 CAPLUS
 CN [1,2,4]Triazolo[1,5-a]pyrazin-8-amine,
 6-(3-cyclopentyl-1-propyn-1-yl)-2-(2-furanyl)- (CA INDEX NAME)



REFERENCE COUNT: 6 THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS
 RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

=> fil stnguide		
COST IN U.S. DOLLARS	SINCE FILE	TOTAL
	ENTRY	SESSION
FULL ESTIMATED COST	11.78	197.88
DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE	TOTAL
	ENTRY	SESSION
CA SUBSCRIBER PRICE	-1.64	-1.64

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COST IN U.S. DOLLARS	SINCE FILE	TOTAL
	ENTRY	SESSION
FULL ESTIMATED COST	0.21	198.09
DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE	TOTAL
	ENTRY	SESSION
CA SUBSCRIBER PRICE	0.00	-1.64

STN INTERNATIONAL LOGOFF AT 15:42:33 ON 07 APR 2009